FEB - 6 2014

Section C

510(k) Summary

"This summary of 510(k) safety and effectiveness information is being submitted in accordance with the requirements of SMDA 1990 and 21 CFR 807.92."

"The assigned 510(k) number is:____K131341____." (applicant leave blank)

Premarket Notification [510(k)] Summary

[(a)(1)]. The summary contains on the first page, preferably on your letterhead paper, the submitter's name, address, phone and fax numbers, name of contact person, and date the summary was prepared :

Submitter's name :	Shijiazhuang Winful Plastic Co., Ltd.
Submitter's address :	No.6 Cangshi Road, Jinzhou City, Hebe i, 052260, China
Phone number :	(86) 31184320503
Fax number :	(86) 31184311294
Name of contact person:	Yin Mingfei
Date the summary was prepared:	January 27, 2014

[(a)(2)]. The name of the device, including the trade or proprietary name if applicable, the common or usual name, and the classification name, if known

Device Name:	Powder Free Yellow Synthetic Vinyl Patient Examination Gloves
Proprietary/Trade name:	Powder Free Yellow Synthetic Vinyl Patient Examination Gloves
Common Name:	Patient examination glove
Classification Name:	Patient examination glove
Device Classification:	I
Regulation Number:	21 CFR 880.6250
Panel:	General Hospital (80)
Product Code:	LYZ

[(a)(3)]. An identification of the legally marketed device to which your firm is claiming substantial equivalence .

Class I* Powder Free Yellow Synthetic Vinyl Patient Examination Gloves that meets all of the requirements of ASTM standard D 5250-06 (Reaffirmation 2011).

Predicate device : POWDER-FREE YELLOW SYNTHETIC VINYL PATIENT EXAMINATION GLOVES, ZHAOYANG PLASTIC CO., LTD k110945

[(a)(4)] A description of the device

Device Description : Powder Free Yellow Synthetic Vinyl Patient Examination Gloves that meets all of the requirements of ASTM standard D 5250-06(Reaffirmation 2011).

-- How the device functions:
PVC films form a barrier to body fluids and bloodborne Pathogens

-- Scientific concepts that form the basis for the device
The PVC rubber is water tight under normal conditions of use. Its tensile properties cause it to conform to the hand, allowing movements necessary for a medical procedure.

-- Physical and performance characteristics such as design, materials and physical properties:
PVC gloves are known to create a barrier to bloodborne pathogens and body fluids. ASTM conforming tensile properties create a glove that is strong and flexible. The glove is manufactured in accordance with the requirements of ASTM D5250 and ASTM D5151 requirements.

[(a)(5)] The summary describes the intended use of the device

Device Intended Use: Powder Free Yellow Synthetic Vinyl Patient Examination Gloves is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.

[(a)(6)] A summary of the technological characteristics of new device compared to the predicate device.
The Powder Free Yellow Synthetic Vinyl Patient Examination Gloves, non sterile are summarized with the following technological characteristics compared to ASTM or equivalent standard.

Features & Description	Predicate Device	Subject Device	Result of Comparison
510(k) Number	K110945	K131341	
Company	ZHAOYANG PLASTIC CO., LTD	Shijiazhuang Winful Plastic Co., Ltd.	--
Product name	POWDER-FREE YELLOW SYNTHETICVINYL PATIENT EXAMINATION GLOVES	Powder Free Yellow Synthetic Vinyl Patient Examination Gloves	--
Product Code	LYZ	LYZ	Same
Size	Small/ Medium/ Large/X large	Small/ Medium/ Large/X large	Small/ Medium/ Large/X large
Intend for use	POWDER-FREE YELLOW SYNTHETIC VINYL PATIENT EXAMINATION GLOVES is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	Powder Free Yellow Synthetic Vinyl Patient Examination Gloves is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	Substantially equivalent
Device Description and Specifications	Meets ASTM D5250-06 (Reapproved 2011)	Meets ASTM D5250 -06 (Reapproved 2011)	Substantially equivalent
Dimensions -- Length	Meets ASTM D5250-06 (Reapproved 2011) ≥230mm min.	230mm min for all sizes	Substantially equivalent
Dimensions -- Width	Meets ASTM D5250-06 (Reapproved 2011) Small 80-90 mmr	Small 80-85 mm Medium 95-97 mm Large 102-108mm	Substantially equivalent

	Medium 90-100mm Large 100-110mm X large 110-120 mm	X large 114-118 mm	
Dimensions -- Thickness	Meets ASTM D5250-06 (Reapproved 2011) Finger 0.05mm min. Palm 0.08mm min.	 Finger 0.05mm min. Palm 0.08mm min.	
Physical Properties	Meets ASTM D5250-06 (Reapproved 2011) Before aging/after aging Elongation ≥300% Tensile Strength≥ I4MPa	Before aging/after aging Elongation ≥300% Tensile Strength≥ I4MPa	Substantially equivalent
Freedom from Pinholes	Meets • 21 CFR 800.20 • ASTM D5250-06 (Reapproved 2011) • ASTM D 5151-06 (Reapproved 2011)	Meets ASTM D5151 Holes Inspection Level I AQL2.5	Substantially equivalent
Residual Powder	Meets ASTM D 6124-06 (Reapproved 2011)	D 6124-06 (Reapproved 2011) Results generated values below 2mg of residual powder	Substantially equivalent
Compare all materials used to fabricate the devices	PVC	PVC	Substantially equivalent
Dusting or Donning Powder:	PU	PU	Substantially equivalent
Dusting or Donning Powder: name	PU	Surface Coating Agent	Substantially equivalent
Compare performance data supporting substantial equivalence	Meets • ASTM D5151-06 (Reapproved 2011) • ASTM D5250-06 (Reapproved 2011) • ASTM D6124-06 (Reapproved 2011)	Meets • ASTM D5151-06 (Reapproved 2011) • ASTM D5250-06 (Reapproved 2011) • ASTM D6124-06 (Reapproved 2011)	Substantially equivalent
Single Patient Use	Single Patient Use	Single Patient Use	Substantially equivalent
Biocompatibility	SKIN IRRITATION DERMAL and SENSITIZATION STUDIES Meets ISO 10993-10	The test article was a non-irritant or non-sensitizer. SKIN IRRITATION DERMAL and SENSITIZATION STUDIES Meets ISO 10993-10	Substantially equivalent
Labeling for the legally marketed device to which substantial equivalence is claimed.	-Powder-free -Patient Examination Glove - Yellow color -non sterile -Single Use Only - Manufactured For: - Lot	-Powder-free -Patient Examination Glove - Yellow color -non sterile -Single Use Only - Manufactured For: - Lot	Substantially equivalent

[(b)(1)] A brief discussion of the non-clinical submitted, reference, or relied on in the premarket notification submission for a determination of substantial equivalence .

Powder Free Yellow Synthetic Vinyl Patient Examination Gloves meet requirements per ASTM D5250-06(Reaffirmation 2011), per ASTM D6124-06(Reaffirmation 2011), per 21 CFR 800.20 and ISO 10993-10: 2002/Amd. 1:2006(E).

[(b)(2)] A brief discussion of the clinical submitted, reference, or relied on in the premarket notification submission for a determination of substantial equivalence.

Clinical data is not needed for gloves or for most devices cleared by the 510(k) process.

[(b)(3)] The conclusions drawn from the non-clinical and clinical tests that demonstrate that the device is as safe, as effective, and performed as well or better than the legally marketed device identified in (a)(3).

It can be concluded that the Powder Free Vinyl Patient Examination Gloves meet the ASTM standard or equivalent standard and FDA requirements for waterleak test on pinhole AQL., meet labeling claims and the Powder Free Yellow Synthetic Vinyl Patient Examination Gloves is as safe, as effective, and performs as well as the predicate device, POWDER-FREE YELLOW SYNTHETIC VINYL PATIENT EXAMINATION GLOVES, ZHAOYANG PLASTIC CO., LTD K110945

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
10903 New Hampshire Avenue
Document Control Center – WO66-G609
Silver Spring, MD 20993-0002

February 6, 2014

Shijiazhuang Winful Plastic Company, Limited
C/O Mr. Chu Xiaoan
Room 1606 Bldg 1 Jianxiang Yuan #209
Bei Si Huan Zhong Road, Haidian District
Beijing 100083
CHINA

Re: K131341
Trade/Device Name: Powder Free Yellow Synthetic Vinyl Examination Gloves
Regulation Number: 21 CFR 880.6250
Regulation Name: Patient Examination Glove
Regulatory Class: I
Product Code: LYZ
Dated: December 19, 2013
Received: December 30, 2013

Dear Mr. Xiaoan:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm.

Sincerely yours,

Erin I. Keith, M.S.
Acting Director
Division of Anesthesiology, General Hospital,
Respiratory, Infection Control and
Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Indications for Use

Form Approved: OMB No. 0910-0120
Expiration Date: December 31, 2013
See PRA Statement on last page.

510(k) Number *(if known)*
K 131341

Device Name
Powder Free Yellow Synthetic Vinyl Patient Examination Gloves

Indications for Use *(Describe)*
Powder Free Yellow Synthetic Vinyl Patient Examination Gloves is a non-sterile disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.

Type of Use *(Select one or both, as applicable)*

☐ Prescription Use (Part 21 CFR 801 Subpart D) ☒ Over-The-Counter Use (21 CFR 801 Subpart C)

PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON A SEPARATE PAGE IF NEEDED.

FOR FDA USE ONLY

Concurrence of Center for Devices and Radiological Health (CDRH) *(Signature)*

Elizabeth F. Claverie -S
2014.02.05 15:36:49 -05'00'

FDA

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